

AMENDMENTS

In the Claims:

1.-10. (Canceled)

11. **(Currently Amended)** A method of inserting an exogenous nucleic acid into the genome of a mouse or rat, said method comprising:

introducing into said mouse or rat a P-element derived vector comprising said exogenous nucleic acid under conditions sufficient for transposition to occur, wherein said P-element derived vector further comprises a pair of P-element transposase recognized insertion sequences flanking ~~a P-foot flanked domain of at least about 2,000 bp in length, wherein said P-foot flanked domain~~ **comprises** a heterologous promoter and a single transcriptionally active gene that comprises said exogenous nucleic acid, wherein said single transcriptionally active gene is separated from one of said P-element transposase recognized insertion sequences by a distance of about 1,000 bp or less, so that said exogenous nucleic acid is inserted into said genome

wherein said P-element derived vector further comprises a transposase domain, or

wherein said method further comprises introducing a second P-element derived vector comprising a transposase domain into said mouse or rat.

12. (Canceled)

13. **(Currently Amended)** The method according to Claim 11, wherein said P-element derived vector comprises a transposase domain.

14. (Currently Amended) The method according to Claim 11 wherein said method further comprises introducing a second vector comprising a transposase domain into said ~~animal~~ mouse or rat.

15. (Previously Presented) The method according to Claim 11, wherein said exogenous nucleic acid ranges in length from about 50 to 150,000 bp.

16.-26. (Canceled)

27. (Currently Amended) A mouse or rat or cells derived from said mouse or rat that ~~has a pair of P-element transposase recognized insertion sequences integrated into the genome of said mouse or rat or cells derived therefrom~~

has/have been transformed with a P-element derived vector comprising a pair of P-element transposase recognized insertion sequences flanking a heterologous promoter and a single transcriptionally active gene that comprises an exogenous nucleic acid.

wherein said single transcriptionally active gene is separated from one of said P-element transposase recognized insertion sequences by a distance of about 1,000 bp or less; and

wherein said P-element derived vector further comprises a transposase domain, or

wherein said mouse or rat or cells has/have been transformed with a second P-element derived vector comprising a transposase domain.

28.-30. (Canceled)

31. (Previously Presented) The composition of claim 27 wherein said mouse or rat or cells derived therefrom has a pair of P-element transposase recognized

31bp insertion sequences integrated into the genome of said mouse or rat or cells derived therefrom.

32.-38. (Canceled)

39. (Previously Presented) The method according to Claim 11, wherein said method is a method of inserting an exogenous nucleic acid into the genome of a mouse.

40. (Previously Presented) The method according to Claim 11, wherein said method is a method of inserting an exogenous nucleic acid into the genome of a rat.

41. (Currently Amended) A method of inserting an exogenous nucleic acid into the genome of a mouse, said method comprising:

introducing into said mouse a P-element derived vector comprising said exogenous nucleic acid under conditions sufficient for transposition to occur, wherein said P-element derived vector comprises a pair of P-element transposase recognized insertion sequences flanking ~~a P-foot flanked domain of at least about 2,000 bp in length, wherein said P-foot flanked domain comprises~~ at least one transcriptionally active gene that is located within 1,000 bp at least 50 bp of one of the P-element transposase recognized sequences; ~~and a transposase domain.~~

wherein said P-element derived vector further comprises a transposase domain, or

wherein said method further comprises introducing a second P-element derived vector comprising a transposase domain into said mouse.

42. (Currently Amended) A method of inserting an exogenous nucleic acid into the genome of a mouse, said method comprising:

introducing into said mouse a P-element derived vector comprising said exogenous nucleic acid under conditions sufficient for transposition to occur, wherein said P-element derived vector comprises a pair of P-element transposase recognized insertion sequences flanking ~~a P-foot flanked domain of at least about 2,000 bp in length, wherein said P-foot flanked domain comprises at least one a heterologous promoter and a single~~ transcriptionally active gene ~~that is within at least 50 bp of one of the P-element transposase recognized sequences,~~

wherein said single transcriptionally active gene is separated from one of said P-element transposase recognized insertion sequences by a distance of about 1,000 bp or less; and

(a) wherein said P-element derived vector further comprises a transposase domain, or

(b) wherein said method further comprises:

(i) inserting a second P-element ~~derived~~ vector comprising a transposase domain into the genome of said mouse; or

(ii) ~~inserting~~ cells derived therefrom.

43. (New) The method according to Claim 41, wherein said P-element derived vector comprises a transposase domain.

44. (New) The method according to Claim 41 wherein said method further comprises introducing a second vector comprising a transposase domain into said mouse.